

MedHealth Review, Inc.

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Notice of Independent Review Decision

DATE NOTICE SENT TO ALL PARTIES: 1/14/15

IRO CASE #:

DESCRIPTION OF THE SERVICE OR SERVICES IN DISPUTE

The item in dispute is the prospective medical necessity of an OP Caudal ESI.

A DESCRIPTION OF THE QUALIFICATIONS FOR EACH PHYSICIAN OR OTHER HEALTH CARE PROVIDER WHO REVIEWED THE DECISION

The reviewer is a Medical Doctor who is board certified in Orthopedic Surgery. The reviewer has been practicing for greater than 10 years.

REVIEW OUTCOME

Upon independent review the reviewer finds that the previous adve	rse
determination/adverse determinations should be:	

Upheld	(Agree)
⊠Overturned	(Disagree)
☐Partially Overturned	(Agree in part/Disagree in part)

The reviewer disagrees with the previous adverse determination regarding the prospective medical necessity of an OP Caudal ESI

A copy of the ODG was not provided by the Carrier or URA for this review.

PATIENT CLINICAL HISTORY [SUMMARY]:

was injured on xx/xx/xx. He sustained low back pain. On 3/25/14, a lumbar MRI revealed a disc protrusion with S1 nerve root abutment. Clinical notes revealed treatment with medications, PT and ESI, including a 60% relief for 6 weeks from the prior ESI injection (as noted on 9/12/14.) Facet injections had also been administered. On 9/12/14, persistent low back pain with 30% right hip pain was

noted. Exam findings included tenderness, positive straight leg raising, and 4+/5 right gastrocsoleus/tibialis anterior strength. Diagnoses included lumbar sprain/strain. On 12/8/14, there was persistent back and right hip pain, paraspinal tenderness and painful lumbar motion. There was also 4+/5 weakness in the right gastrocsoleus and tibialis anterior muscles. Denial letters included the lack of adequate guideline-associated response from the prior injection.

ANALYSIS AND EXPLANATION OF THE DECISION INCLUDE CLINICAL BASIS, FINDINGS AND CONCLUSIONS USED TO SUPPORT THE DECISION.

The claimant does appear to have objective evidence of radiculopathy. The prior epidural steroid injection resulted in six weeks of a documented 60% relief of the overall symptoms. Therefore, applicable clinical guidelines have been fully met (as referenced below) with regards to a consideration for a repeat epidural steroid injection. The request is considered medically reasonable and necessary at this time.

Reference: Low Back Chapter Criteria for the use of Epidural steroid injections: Note: The purpose of ESI is to reduce pain and inflammation, thereby facilitating progress in more active treatment programs, reduction of medication use and avoiding surgery, but this treatment alone offers no significant long-term functional benefit.

- (1) Radiculopathy (due to herniated nucleus pulposus, but not spinal stenosis) must be documented. Objective findings on examination need to be present. Radiculopathy must be corroborated by imaging studies and/or electrodiagnostic testing.
- (2) Initially unresponsive to conservative treatment (exercises, physical methods, NSAIDs and muscle relaxants).
- (3) Injections should be performed using fluoroscopy (live x-ray) and injection of contrast for guidance.
- (4) Diagnostic Phase: At the time of initial use of an ESI (formally referred to as the "diagnostic phase" as initial injections indicate whether success will be obtained with this treatment intervention), a maximum of one to two injections should be performed. A repeat block is not recommended if there is inadequate response to the first block (< 30% is a standard placebo response). A second block is also not indicated if the first block is accurately placed unless: (a) there is a question of the pain generator; (b) there was possibility of inaccurate placement; or (c) there is evidence of multilevel pathology. In these cases a different level or approach might be proposed. There should be an interval of at least one to two weeks between injections.
- (5) No more than two nerve root levels should be injected using transforaminal blocks.
- (6) No more than one interlaminar level should be injected at one session.
- (7) Therapeutic phase: If after the initial block/blocks are given (see "Diagnostic Phase" above) and found to produce pain relief of at least 50-70% pain relief for at least 6-8 weeks, additional blocks may be supported. This is generally referred

to as the "therapeutic phase." Indications for repeat blocks include acute exacerbation of pain, or new onset of radicular symptoms. The general consensus recommendation is for no more than 4 blocks per region per year.

- (8) Repeat injections should be based on continued objective documented pain relief, decreased need for pain medications, and functional response.
- (9) Current research does not support a routine use of a "series-of-three" injections in either the diagnostic or therapeutic phase. We recommend no more than 2 ESI injections for the initial phase and rarely more than 2 for therapeutic treatment.
- (10) It is currently not recommended to perform epidural blocks on the same day of treatment as facet blocks or sacroiliac blocks or lumbar sympathetic blocks or trigger point injections as this may lead to improper diagnosis or unnecessary treatment.
- (11) Cervical and lumbar epidural steroid injection should not be performed on the same day. (Doing both injections on the same day could result in an excessive dose of steroids, which can be dangerous, and not worth the risk for a treatment that has no long-term benefit.)

A DESCRIPTION AND THE SOURCE	OF THE SCREENING CRITERIA OR
OTHER CLINICAL BASIS USED TO I	MAKE THE DECISION:
☐ ACOEM- AMERICAN COLLEG	E OF OCCUPATIONAL &
ENVIRONMENTAL MEDICINE	UM KNOWLEDGEBASE
☐ AHCPR- AGENCY FOR HEAL	THCARE RESEARCH & QUALITY
GUIDELINES	
☐ DWC- DIVISION OF WORKER	S COMPENSATION POLICIES OR
GUIDELINES	
EUROPEAN GUIDELINES FOR	R MANAGEMENT OF CHRONIC LOW
BACK PAIN	
☐ INTERQUAL CRITERIA	
	ICAL EXPERIENCE AND EXPERTISE IN
ACCORDANCE WITH ACCEP	TED MEDICAL STANDARDS
	S CONFERENCE GUIDELINES
☐ MILLIMAN CARE GUIDELINES	3
ODG- OFFICIAL DISABILITY (SUIDELINES & TREATMENT
GUIDELINES	
PRESSLEY REED, THE MEDIC	
☐ TEXAS GUIDELINES FOR CHI	ROPRACTIC QUALITY ASSURANCE &
PRACTICE PARAMETERS	
☐ TEXAS TACADA GUIDELINES	;
☐ TMF SCREENING CRITERIA IN	//ANUAL
	LY ACCEPTED MEDICAL LITERATURE
(PROVIDE A DESCRIPTION)	
 ·	CIENTIFICALLY VALID, OUTCOME
FOCUSED GUIDELINES (PRO	VIDE A DESCRIPTION)